



Product Service

EC - CERTIFICATE

Product Quality Assurance System

(Annex VI of the Directive 93/42/EEC on Medical Devices)

No. G3M 10 09 20267 015

Manufacturer: Friedrich Bosch GmbH & Co. KG

Hohenlaidenstr. 30
72406 Bisingen
GERMANY

Facility(ies):

Friedrich Bosch GmbH & Co. KG
Hohenlaidenstr. 30, 72406 Bisingen, GERMANY

Product Category(ies): Mechanical Sphygmomanometers and Pressure Infusion Cuffs

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for final inspection and test according to Annex VI, section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system covers those aspects concerned with the conformity with the metrological requirements of the respective product / product categories and complies with the provisions of this Directive. It is subject to periodical surveillance. See also notes overleaf.

Report no.: 71375338

Valid until: 2015-11-15

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Hans-Heiner Junker



TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.

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